

### **Amendments to the Claims**

This listing of claims replaces all prior versions and listings of claims in the application. Any amendments or cancellations to the claims are made without prejudice or disclaimer.

#### **1-44. (Canceled)**

45. **(Previously Presented)** A substantially pure consecutive and anti-angiogenic polypeptide, comprising the central region of human histidine rich glycoprotein (HRGP) corresponding to SEQ.ID.NO:2.

46. **(Previously Presented)** The polypeptide of claim 45, consisting of the central region of human histidine rich glycoprotein (HRGP) corresponding to SEQ.ID.NO:2.

47. **(Previously Presented)** A substantially pure consecutive and anti-angiogenic polypeptide, consisting of a subfragment of the central region of human HRGP (SEQ.ID.NO:2).

48. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to region 330-364 (SEQ.ID.NO: 1) of mature human HRGP.

49. **(Previously Presented)** The polypeptide of claim 47, said subfragment having an amino acid length of between 3 and 35 amino acids.

50. **(Previously Presented)** The polypeptide of claim 49, having an amino acid length selected from the group consisting of between 3 and 25 amino acids, 3 and 20 amino acids, 3 and 15 amino acids, 3 and 10 amino acids, and 3 and 8 amino acids.

51. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO: 18.

52. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO: 17.

53. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO: 16.

54. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:22.

55. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:21.

56. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:24.

57. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:23.

58. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:26.

59. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:25.

60. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:28.

61. **(Previously Presented)** The polypeptide of claim 47, said subfragment corresponding to SEQ.ID.NO:27.

62. **(Previously Presented)** The polypeptide of claim 47, wherein said polypeptide is isolated from human HRGP.

63. **(Previously Presented)** The polypeptide of claim 47, wherein said polypeptide is isolated from proteolytically processed human HRGP purified from plasma.

64. **(Previously Presented)** The polypeptide of claim 47, wherein said polypeptide is recombinantly produced or isolated from recombinantly produced human HRGP.

65. **(Previously Presented)** The polypeptide of claim 47, wherein said polypeptide is synthetically produced.

66. **(Previously Presented)** The polypeptide of claims 47, wherein said polypeptide does not promote angiogenesis or does not bind to thrombospondin.

67. **(Previously Presented)** A pharmaceutical composition comprising an effective amount of the polypeptide of claim 47.

68. **(Previously Presented)** The pharmaceutical composition of claim 67, further comprising a pharmaceutically acceptable carrier.

69. **(Previously Presented)** The pharmaceutical composition of claim 67, further comprising an anti-angiogenic agent.

70. **(Previously Presented)** The pharmaceutical composition of claim 69, wherein said anti-angiogenic agent is selected from the group consisting of angiostatin, thrombostatin, endostatin, interferon- $\alpha$ , interferon-inducible factor 10, and platelet factor 4.

71. **(Previously Presented)** The pharmaceutical composition of claim 67, further comprising an anti-neoplastic agent.

72. **(Previously Presented)** The pharmaceutical composition of claim 71, wherein said antineoplastic agent is selected from the group consisting of taxol, cyclophosphamide, carboplatinum, cisplatin, gancyclovir, camptothecin, paclitaxel, hydroxyurea, 5-azacytidine, 5-aza-2'-deoxycytidine, and suramin.

73. **(Previously Presented)** The pharmaceutical composition of claim 67, further comprising an anti-inflammatory agent.

74. **(Previously Presented)** The pharmaceutical composition of claim 73, wherein said antiinflammatory agent is selected from the group consisting of prednisone, a cox-2 inhibitor, ibuprofen and aspirin.

75. **(Previously Presented)** The pharmaceutical composition of claim 67, further comprising an effective amount of  $Zn^{2+}$ .

76-83. **(Canceled)**